

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
15 September 2005 (15.09.2005)

PCT

(10) International Publication Number
WO 2005/085313 A1

(51) International Patent Classification⁷: C08G 18/10,
C09D 167/07, 175/16, C09J 167/07, 175/16

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(21) International Application Number:
PCT/AU2005/000306

(81) Designated States (*unless otherwise indicated, for every
kind of national protection available*): AE, AG, AL, AM,
AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN,
CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE,
KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,
MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG,
PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ,
TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA,
ZM, ZW.

(22) International Filing Date: 3 March 2005 (03.03.2005)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
2004901109 3 March 2004 (03.03.2004) AU

(84) Designated States (*unless otherwise indicated, for every
kind of regional protection available*): ARIPO (BW, GH,
GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM,
ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI,
FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO,
SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN,
GQ, GW, ML, MR, NE, SN, TD, TG).

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Published:

— with international search report

*For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.*

(54) Title: POLYMER COMPOSITIONS FOR DUAL OR MULTI STAGED CURING

(57) Abstract: The present invention provides a polymer composition for use at least in surface coating applications and in fabrication of rigid foams with load bearing capacity comprising a base molecule, a linker molecule and at least one initiator compound, said base molecule having at least two differing functionalities, and said linker molecule having a functionality reactive with at least one functionality of said base molecule, the first of said at least two functionalities of said base molecule enabling a first curing stage of said polymer composition by reaction with said linker molecule, and the second and any further functionality of said base molecule enabling second and further curing stages of said polymer composition, said first, second and any further curing stages being capable of activation simultaneously or independently of each other as required. A further provides uses of the polymer compositions of the invention in surface coating and adhesives applications and in the preparation of rigid foams with load bearing capacity.



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POLYMER COMPOSITIONS FOR DUAL OR MULTI STAGED CURING

FIELD OF INVENTION

The present invention relates to polymer compositions for use in surface coating such as in antifouling and antibacterial applications, as adhesives, sealants, as binders in inks and in lamination processes and in forming porous polymer constructs. The polymers are capable of curing on demand in two or more stages which are of the same or different character, and which are carried out simultaneously, in tandem or step wise. The compositions may be degradable or stable. During the first stage of curing the polymer may be foamed to construct the porous structure with desired pore morphologies and mechanical properties.

BACKGROUND

The nature and chemistry of synthetic polymers used in surface coating, in adhesive, in sealant and in printing applications varies depending on the application. In general they need to have the advantage over other materials of being able to be tailored as far as their mechanical properties are concerned to the application at hand. Such polymers can also be tailored to contain various functionalities allowing them to integrate into the environment in which they are to be placed.

A vast majority of biodegradable polymers studied belong to the polyester family. Among these poly(α -hydroxy acids) such as poly(glycolic acid), poly(lactic acid) and a range of their copolymers have historically comprised the bulk of published material on biodegradable polyesters and have a long history of use as synthetic biodegradable materials in a number of applications. Among these applications, poly(glycolic acid), poly(lactic acid) and their copolymers, poly(p-dioxanone), and copolymers of trimethylene carbonate and glycolide have been the most widely used.

Almost all of these polymers, both biodegradable and biostable, are pre-manufactured or pre-cured and moulded, prior to application. In a minority of cases two or many different reactive species are mixed together immediately prior to application, so that the user has a specified window of time before the polymer begins to set or cure and becomes unworkable.

Polymer compositions that can cure by multiple curing methods and steps which are designed for applications such as surface coatings and adhesives are reported in the literature. The technique is widely used in surface coating and in adhesive formulations. In almost all cases, thermal and radiation (UV or other
5 sources of radiation) have been the most widely used curing methods. Typically the coating compositions consist of a mixture of compounds with appropriate functional groups that can react by thermal or radiation initiated free radical mechanisms. One example is described in US application 2003083397 which
10 discloses dual cure coating compositions containing a polyester, acrylated aliphatic urethane oligomer, polyester tackifier processing additive, pigment, catalyst, photoinitiator and a triisocyanate. Such mixtures provide good surface coatings but are unnecessarily complex and are not easily manipulated.

Other compositions described in the literature suffer issues with their durability, and their rheology during application. Polyisocyanate based
15 compositions are known as being insufficiently viscous at the high temperatures required for curing.

It is one object of this invention to provide multifunctional polymer compositions for use at least in surface coating applications, the curing process of which may be controlled to allow ease of application, and the design of which
20 may be adjusted to suit the specific application. The polymers are desirably capable of incorporating components such as pigment which maybe an adjunct in the application selected.

It is another object of this invention to provide multifunctional polymer compositions for use in the fabrication of rigid foams, particularly those with load
25 bearing capacity. The multi-stage curing feature of the compositions in this invention allows one to use the first stage curing to build the foam/construct with desired pore density/size etc while the polymer composition is in a mouldable state and carry out subsequent curing steps to achieve full cure and the high strength.

30 SUMMARY OF INVENTION

To this end there is provided a polymer composition for use at least in surface coating and adhesive applications and in fabrication of rigid foams with load bearing capacity comprising a base molecule, a linker molecule and at least one initiator compound, said base molecule having at least two differing

functionalities, and said linker molecule having a functionality reactive with at least one of said functionalities of said base molecule, the first of said at least two functionalities of said base molecule enabling a first curing stage of said polymer composition by reaction with the functionality of said linker molecule, and the
5 second and any further functionality of said base molecule enabling second and optionally further curing stages of said polymer composition, said first, second and any further curing stages being capable of activation simultaneously or independently of each other as required.

This invention also provides a prepolymer composition for use at least in
10 surface coating and adhesive applications and in the fabrication of rigid foams with load bearing capacity comprising the reaction product of a base molecule, a linker molecule and at least one initiator compound, said base molecule having at least two differing functionalities, and said linker molecule having a functionality reactive with at least one of said functionalities of said base molecule, said
15 reaction product being the result of a first curing stage wherein the first of said at least two functionalities of said base molecule reacts with the functionality of said linker molecule to form said prepolymer composition.

Preferably the prepolymer composition is flowable and injectable.

Throughout this specification the term "prepolymer" will be used to mean
20 the polymer composition which is formed in a first stage of curing by combination of a base molecule and a linker molecule but which is yet to be further cured in a second or further curing stage.

This invention also provides a cured polymeric end product for use at least in surface coating and adhesive applications and in the fabrication of rigid foams
25 with load bearing capacity comprising the reaction product of a base molecule, a linker molecule and at least one initiator compound, said base molecule having at least two differing functionalities, and said linker molecule having a functionality reactive with at least one of said functionalities of said base molecule, said end product being the result of a first curing stage wherein the first of said at least two
30 functionalities of said base molecule reacts with the functionality of said linker molecule and a second curing stage and optionally further curing stages wherein said initiator compound is activated to affect free radical polymerisation of at least said second functionality of said base molecule.

It has been found that the compositions according to the invention are particularly useful in applications where two or more independent and non-interfering curing methods are desired. The first mode of curing might for example be used to increase the viscosity of the composition such that the material could be shaped or moulded to desired shapes and forms, and the second mode of curing might be used to fix the final shape. The compositions according to the invention may also be engineered to cure in the presence of selective functional groups, causing strong covalent bonding to the substrate, or cured to a hard, space filling material by cross linking either simultaneously and/or in tandem and/or step wise as required. The polymers have the option of being engineered to be biodegradable or biostable for permanent fixtures and cured on demand in either aqueous or organic environments. They can be made flowable. When water is present in the compositions, the end cured material is porous enabling the delivery of other components suited to the application. It has been found that the compositions of the invention may be typically cured on demand to a hard, optionally porous material with 1-2 minutes of irradiation.

Further, the multi-stage curing feature of the compositions according to the present invention is particularly suitable for fabricating foams with closed or interconnected pores. For example, a prepolymer with isocyanate and olefinic functional groups could first be reacted with a mixture of water and surfactants to increase viscosity and to generate pores with release of carbon dioxide by the reaction of isocyanate groups and water. The composition at this stage would be either formed to a desired shape or size and then cured in a mould to the desired shape using subsequent curing methods. The choice of linker and base molecules allows one to prepare such foamed structure with high compressive strength as well as with high load bearing capacity. Second curing stages may be carried out either through thermal or other means of initiation to polymerize the olefinic moieties in the prepolymer.

Preferably the base molecule and the linker molecule may each be, independently of the other, a single molecule, or alternatively an oligomer formed from two or more substrate monomers. When the base or linker molecule is oligomeric, it has a molecular weight of less than 2000, preferably less than 1000 and more preferably less than 500. The compositions of the invention may also comprise a radical inhibitor to prevent premature polymerisation of a prepolymer

composition during storage, and/or a sensitizer or a promoter to assist in the activation of said second and/or further curing steps.

Still more preferably the compositions of the invention may also comprise a dispersant or porogen such as for example, but not limited to water; or other
5 particulate components.

As a porogen, water is preferably present in the compositions of the invention in an amount of up to 40% of the total weight of the composition. Higher levels of water may be incorporated if an emulsifier is also present in the composition. In such instances water levels in an amount of up to 80% of the
10 total weight of the composition may be incorporated. Addition of an emulsifier may also help to control pore size and distribution. Any emulsifier could be used but emulsifiers such as block copolymers of polyethylene glycol and polypropylene glycol (Pluronic available from BASF), block copolymers of polysiloxane and polyethylene glycol are preferred for biomedical applications.
15 Commercially available emulsifiers that may be suitable include Symperonic PEF127 and Symperonic PE L101 (Uniqema).

Throughout this specification the term "initiator compound" shall be taken to mean a molecule or mixture of molecules which when activated by an energy source, will result in free radical polymerization of said composition in a second
20 and/or further curing step. The second and/or further curing step is preferably activated by means external of the composition such as, for example, irradiation by light of particular wavelength (photoinitiation), by thermal initiation, or by redox initiation with the result that free radical polymerization occurs so as to harden the polymer composition. The terms "dual cure", "multiple cure" and "cure on
25 demand" are interchangeably used to indicate a polymer that may be cured in more than one stage and at the timing of the users choice by application of suitable initiator stimulus.

In photoinitiated polymerisations, a sensitizer may be added to increase the rate of initiation. Other organic additives may be incorporated to shift the
30 wavelength at which polymerisation occurs. The sensitizer may be a separate molecule which is of a chemical nature to predispose said polymer composition to a second or subsequent curing step under certain environmental conditions. Thus throughout this specification, the term "sensitizer" shall be taken to mean any

molecule which assists in the activation and increases the rate of photoinitiation of a second or subsequent curing step.

Similarly, in polymerisations initiated thermally, a 'promoter' may be added to accelerate the rate of initiation and/or to initiate polymerisation at a desired

5 temper

A catalyst may also be added to the compositions of the invention to assist in the curing processes. Surfactants may also be added so as to modify the characteristics of the resultant process.

The invention also provides a use of the prepolymer, polymer and cured
10 end products of the invention in the preparation of a surface coating.

The invention also provides a use of the prepolymer, polymer and cured end products of the invention in the preparation of a rigid foam with load bearing capacity.

The invention also provides a use of the prepolymer, polymer and cured
15 end products of the invention in the preparation of an adhesive composition.

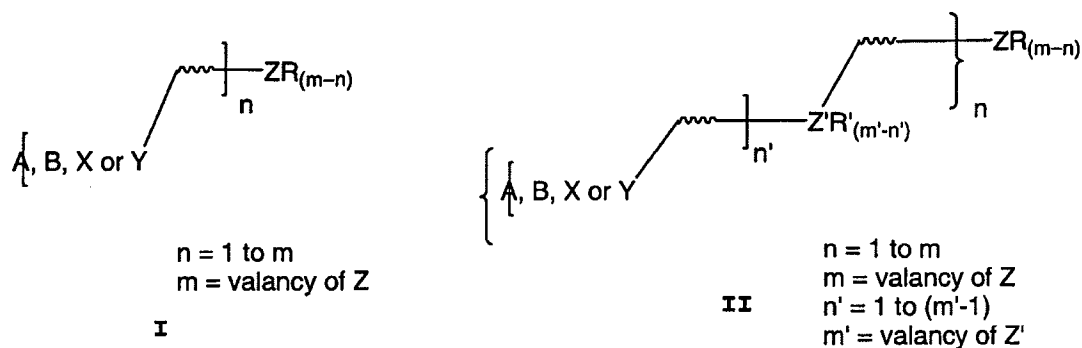
Throughout this specification the terms "comprises" or "comprising" or grammatical variations thereon shall be taken to specify the presence of stated features, integers, steps or components but does not preclude the presence or addition of one or more other features, integers, steps, components or groups
20 thereof not specifically mentioned.

DETAILED DESCRIPTION OF THE INVENTION

In the compositions according to the invention the base molecule and linker molecule are preferably the two major components by mass and may, in general, either or both be a single compound or an oligomer of appropriately low
25 molecular weight which when mixed and/or reacted produce a prepolymer composition that is able to be shaped, is preferably flowable and is not rendered solid until such time as a second or further curing step is activated by an external method. The base and linker molecules desirably each may possess at least one functional group that is able to bond to a functional group of the other molecule,
30 preferably using an ionic bond, and more preferably a covalent bond that has sufficient bonding characteristics as to maintain the composition in the desired physical form in the environment in which it is placed until such time as a second or further curing step is affected. Similarly, the initiator, sensitizer and promoter molecule may contain such functional groups. However, it is not essential that the

initiator, sensitizer or promoter molecules have such properties for the compositions to have cure on demand characteristics.

Preferred general structure of the base is represented by the general formulae I or II.



segment may be one or more repeating units. n, m, n' and m' are integers.

5

In the base molecule designated by formula I, Z may be H, C, O, N, Si, S. The remaining variables may then be defined according to the following table:

Table 1:

Atom (Z)	Valancy (m)	n (arms)	R	ZR(m-n)
C	4	1	H, CH ₃ , C ₂ H ₅ or OR	CH ₃ , C(CH ₃) ₃ , C(C ₂ H ₅) ₃ , C(OR) ₃
		2	H, CH ₃ , C ₂ H ₅ or OR	CH ₂ , C(CH ₃) ₂ , C(C ₂ H ₅) ₂ , C(OR) ₂
		3	H, CH ₃ , C ₂ H ₅ or OR	CH, C(CH ₃), C(C ₂ H ₅), C(OR)
		4	none	C (4 arms)
O	2	1	H, CH ₃ , C ₂ H ₅ or OR	OH, O(CH ₃), O(C ₂ H ₅), R cannot be OR
		2	none	O
Si	4	1	H, CH ₃ , C ₂ H ₅ or OR	SiH ₃ , Si(CH ₃) ₃ , Si(C ₂ H ₅) ₃ , Si(OR) ₃
		2	H, CH ₃ , C ₂ H ₅ or OR	SiH ₂ , Si(CH ₃) ₂ , Si(C ₂ H ₅) ₂ , Si(OR) ₂
		3	H, CH ₃ , C ₂ H ₅ or OR	SiH, Si(CH ₃), Si(C ₂ H ₅), Si(OR)
		4	none	Si (4 arms)
H	1	1	none	H (R=0)

10 In the linker molecule designated by formula II, Z' may be H, C, O, N, Si, S. The remaining variables are as defined in Table 1 and in the following table:

Table 2:

Atom (Z')	Valancy (m')	n' (arms)	R'	Z'R'(m'-n')
C	4	1	H, CH ₃ , C ₂ H ₅ or OR	CH ₂ , C(CH ₃) ₂ , C(C ₂ H ₅) ₂ , C(OR) ₂
		2	H, CH ₃ , C ₂ H ₅ or OR	CH, C(CH ₃), C(C ₂ H ₅), C(OR)
		3	none	C (4 arms)
O	2	1	none	O
Si	4	1	H, CH ₃ , C ₂ H ₅ or OR	SiH ₂ , Si(CH ₃) ₂ , Si(C ₂ H ₅) ₂ , Si(OR) ₂
		2	H, CH ₃ , C ₂ H ₅ or OR	SiH, Si(CH ₃), Si(C ₂ H ₅), Si(OR)
		3	none	Si (4 arms)
H	1	0	cannot have a group or atom	

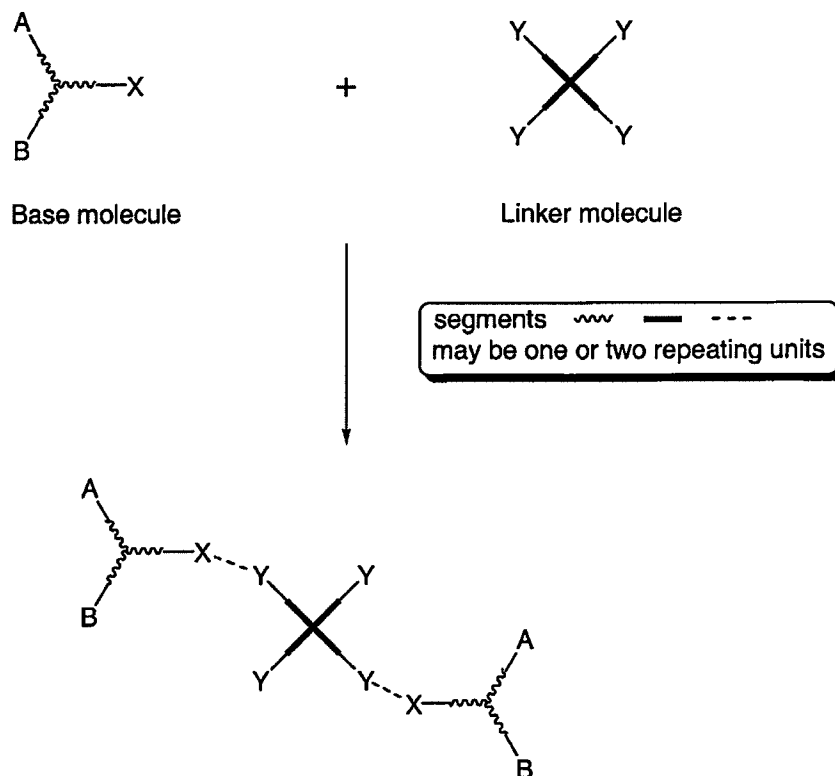
Functional groups A and B may be unsaturated moieties that are independently radically polymerizable depending on the method and type of initiator used. Functional group A is preferably different to that of B in the way it may be triggered to polymerize. Functional group A for example may be an acrylate double bond which can be initiated to propagate polymerization by photolytic initiators and/or thermal initiators. Functional group B may for example, be an allyl moiety which can only be initiated under specific free radical initiating conditions, but not by photochemical means. Alternatively the functional group B may only be triggered to polymerise by a different combination of initiators.

The functional group B may not be an ethylenically unsaturated moiety. However, it may be any other moiety that can be triggered to initiate polymerization of unsaturated double bonds.

A linker molecule is a single organic molecule or alternatively an oligomer formed from two or more substrate molecules. The functional groups in the linker molecule may be chosen depending on the X or Y functional groups in the base molecule. For examples, if the base molecule is isocyanato ethylmethacrylate, then the linker molecule can be any polyol or polyamine. Examples include di or higher hydroxy or amine functional oligomers based on polyesters, polycarbonates, polyethers, polysiloxanes etc. Similarly, if the X or Y functional group in the base molecule is hydroxy or amine, then the linker molecule may be

designed to have isocyanate functional groups by reacting said polyols with a diisocyanate.

A typical reaction scheme between a base and a linker molecule is thus as follows:



A: polymerizable by photolytic or thermal activation
 B: polymerizable by thermal activation only
 X-Y: reactive groups at ambient or elevated temperatures
 X or Y: may be used to generate voids leading to porosity
 X or Y may be used to bond to or compatibilise with the environment

5

In the above reaction scheme, the base molecule contains A, B and X functional groups, the examples of which are acrylic, allylic and hydroxyl, respectively. The linker molecule contains Y functional group, examples of which are isocyanate and thioisocyanate. The base and linker molecules are covalently linked via the reaction of X and Y functional groups.

The functional group X (or Y) in the base molecule or the linker molecule may be used for a number of purposes. Firstly, it may be used to bond, either by covalent or ionic means, to the linker. For example, X (or Y) may be hydroxy, isocyanate, carboxy, halo or any other suitable functional group. In such cases the functional group Y (or X) in the linker respectively may be isocyanate, hydroxy

15

or mercapto, halo, carboxy or any other suitable functionality. More particularly, it is desirable to have groups that are highly reactive with each other, such as isocyanate with hydroxyl, amino or thiol functionalities, so as to achieve consistency of the prepolymer composition before application to the environment of use. Secondly, the functional group may be used to enhance integration with the surrounding environment, by skilfully manipulating the mass of residual functionality following the reaction. For example, when X is hydroxy, the remaining groups can promote mixing and enhance greater integration with aqueous surroundings or water as a component in the composition, through hydrogen bonding. Thus there is scope for the number of residual groups present to be varied depending on the application. Thirdly, the group X may be used to generate porosity if desired. For example isocyanate on reaction with water produces carbon dioxide, which may be used as a blowing agent to create voids and/or channel in the polymer immediately prior to curing to obtain a porous polymer where necessary.

The base and linker molecules can be either premixed to afford a one component curable prepolymer system or formulated immediately prior to the application.

The architecture of the prepolymer formed by polymerisation in a first stage of said base and linker molecules may be linear or regularly branched such as for example a star polymer or dendritic or may be irregularly branched such as for example, hyperbranched.

The catalyst may be selected from, but is not limited to tin catalysts such as stannous (II) ethylhexanoate, stannous oleate, stannous chloride, dibutyltin dilaurate (DBTDL), dibutyltin dioxide, dibutyltin di-2-ethylhexanoate; tertiary amine catalysts such as triethylene diamine, tetramethylbutanediamine (TMBDA), dimethylethanolamine, 1,8-diazabicyclo[5.4.0]undec-7-ene, 1,4-diazo[2,2,2]bicycle-octane (DABCO), hydroxyguanine, tetramethyl guanidine, N-ethylmorpholine, riboflavin; titanium catalysts such as titanium ethanol amine, Tyzor-titanate (Tyzor 131), Tyzor organotitanates, titanium butoxide; titanium aqueous chelates which are stable in water such as Tyzor-LA (aqueous titanium lactate chelate), Tyzor 131 (aqueous titanium chelate), Tyzor 217 (aqueous zirconium lactate), Tyzor 218(aqueous zirconium glycolate); and other catalysts such as calcium phosphate, ovalbumin, sodium acetate, and tributyl phosphine.

The selection of the initiator present in the composition for the purpose of triggering free radical curing, is dependant on the method of initiation selected. Initiation may be thermal, photolytic or based on a redox system of components and is preferably by an external source. For example, camphorquinone, phosphine oxide based initiators such as 2,4,6- trimethyl benzoyl) diphenyl phosphine oxide are suitable and redox initiators such as ammonium persulfate and sodium metabisulfite are also suitable. Preferable is a system that cures the polymeric composition using a wave length that is either in the UV or visible region of electromagnetic radiation although actinic radiation may also be used.

In one embodiment of the invention, visible light source having a maximum wave length of $450 \pm 30\text{nm}$ is used. Examples of photoinitiators include but are not limited to 2,2-dimethoxy-2-phenylacetophenone (Irgacure 651), Hydroxyalkyl phenones (1-hydroxycyclohexyl phenyl ketone (Irgacure 184), 2-methyl-1-[4-(methylthio)phenyl]-2-(4-morpholinyl)-1-propanone (Irgacure 907), 2-hydroxy-1-[4-(hydroxyethoxy)phenyl]-2-methyl-1-propanone (Darocur 2959), Darocur 4265, Darocur TPO, Darocur 1173, Irgacure 500, 784, 907, 2959, 819, 2020, 1000, 369, 651, 1300, 819/819W, Irgacure 2005 and Irgacure 2010W and Irgacure 2020, polysilanes, Esacure KP150 (Hydroxyalkylphenylketone), Camphorquinone, Rosebengal, ethyl-4-N,N-dimethylaminobenzoate(4EDMAB)/ triethanolamine, α alkoxydeoxybenzoins, α,α -dialkoxyacetophenone (DEAP), (1-hydroxycyclohexylphenylketone), dibenzoyl disulphide, S-phenyl thiobenzoates, acylphosphine oxide, dibenzoylmethanes, O-acyl α - oximinoketones, phenylazo-4-diphenyl sulphone, benzophenones, flourenones, xanthenes, thioxanthenes, benzils, ketals (2,2-dimethoxy-2-phenylacetophenone DMP), α -ketocoumarines, anthraquinone, and terephthalophenones.

2,6-di-tert-butyl-4-methylphenol may be included in the composition as a radical inhibitor, and helps to avoid premature second stage polymerisation during storage of the prepolymer composition.

Dispersants, such as water, may be used in the composition and with the aim of generating or tailoring the desired physical properties of the resultant cured polymer during the curing process. For example, use of water as a dispersant in a composition that contains residual isocyanate groups will generate carbon dioxide during curing while the other mechanism of curing cross links the material into a hard material. Such dual mode curing enables the generation of voids

and/or channels within the polymer enabling the fabrication of a porous polymer. Alternatively dispersants that do not generate carbon dioxide can be added if porosity is undesirable.

Sensitizers may include but are not limited to bis-(N,N tetraethyl) ethylene
5 diamine, N,N'-dimethyl amino ethyl methacrylate, Ethyl-4-dimethylaminobenzoate, Isopropyl thioxanthone (Quantacure ITX), Ethyl-p-diaminobenzoate, Triethanolamine, Tertiary amine (N,N-diethylaminomethacrylate), and Michler's ketone.

Variation of each component in the composition may be used to dictate the
10 chemical and physical characteristics of the final cured polymer composition. For example, lowering the percentage of acrylate groups has the benefit of tailoring the final material to be softer while an increase makes it otherwise. This may be achieved by adding compounds incorporating acrylate groups in excess in the preparation of the prepolymer. Thus, the desired mechanical properties of the
15 materials can be tailored to the application at hand. Increasing the quantity of dispersant for example lowers the viscosity of the compositions.

Preferred compositions are those in which the ratio of NCO:acrylate functionalities is in the range of 1.0-3.0: 0.5-3.0.

Variation of the initiator concentrations may control the time frame in which
20 the polymer can be cured in to a soft or a hard material and also has an effect on the curing mechanisms. For example, fast curing (with high concentration of initiator) limits the swelling of the polymer composition during the generation of the porosity by carbon dioxide evolution in a dual mode isocyanate/water curing and radical cross linking system.

25 The compositions of the invention may be engineered to cure in either an aqueous or an organic environment and to have an injectable viscosity or to be formed as a solid to suit the application at hand.

In one embodiment disclosed herein the base molecule is monomeric or oligomeric, and can be cured upon incorporation into the prepolymer by free
30 radical polymerization. It may possess one or many isocyanate reactive functional groups, such as but not limited to amino, hydroxyl and thiol and suitably activated double bond groups. More preferably, a monomeric base molecule possessing one or more free radically polymerizable acrylate functional groups and one

hydroxy group reactive towards isocyanates, such as glycerol dimethylacrylate is used.

In an alternative embodiment herein disclosed the base molecule is monomeric or oligomeric containing acrylate and isocyanate functionalities. For
5 example, the base molecule isocyanatoethylmethacrylate (IEM) is reacted with a linker molecule having hydroxy or amine functional groups to produce a prepolymer composition which may be cured in a second stage.

In one preferred embodiment the composition preferably comprises a base molecule which is a polyurethane/urea structure which is more preferably
10 the reaction product of a core molecule with two or more functional groups, either linear or multi arm such as but not limited to degradable polyesters, and an organic molecule containing isocyanate groups, more preferably a diisocyanate. The linker molecule is then reacted with the base molecule such that the resulting prepolymer contains terminal isocyanate end groups as well as free
15 radically polymerizable olefinic groups such as acrylates so as to engineer the chemical and physical characteristics of the desired material.

The prepolymer compositions that are preferred for preparing biodegradable cured end-products are based on degradable polyesters, such as polycaprolactone diol, triol, poly(lactic acid) diol, triol, poly(glycolic acid) diol triol,
20 and copolymer diols and triols of lactic and glycolic acids. A range of architectures of these polyols are commercially available and vary from linear, branched to star type and their suitability to a particular application is solely depended on the requirements of the application at hand. Other polyols could be synthesized based on literature reported procedures.

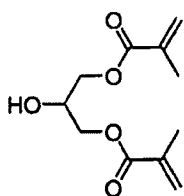
25 For biostable compositions, preferred oligomers for use as linker molecules are prepared by reacting polyether polyols, polysiloxane polyols, hydrocarbon polyols or mixtures thereof with diisocyanates such as 4,4-methylene diphenyl diisocyanate (MDI). Examples of polyether polyols include poly(teramethylene oxide), poly(hexamethylene oxide), poly(octamethylene oxide), and
30 poly(decamethylene oxide). Siloxane based macrodiols such as α , ω -bis(6-hydroxy-ethoxypropyl)-polydimethylsiloxane (available from Shin-Etsu, Japan) are particularly preferred for biostable materials.

Preferred polymer compositions are those based on combinations of one or more of polycaprolactone diol (400-2000), polycaprolactone triol, poly(lactic

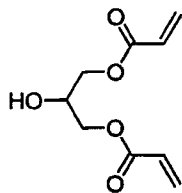
acid) diol, polytetraemthylene ether glycol, glycerol with one or more of ethyl 2,6-diisocyanato hexanoate (ethyl lysinediisocyanate), 4,4-methylene bis(phenyl isocyanate), methy 2,6-diisocyanato hexanoate (methyl lysinediisocyanate), hexane diisocyanate, butane diisocyanate The olefinic functionality is introduced
5 by the use of one or more of isocyanato methacrylate, polyethylene glycol acrylate, glycerol dimethacrylate or isocyanato ethyl methacrylate.

Particularly suited linker molecules may be prepared by reacting di- or higher- hydroxy or amine functional oligomers based on polyesters, polycarbonates, polyethers, polysiloxanes etc with a diisocyanate or
10 polyisocyanate. For biostable polymer compositions polysiloxane, polycarbonate and polyether based polyols are preferred.

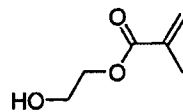
Particularly suited base molecules may be selected from the group consisting of, but not limited to the following:



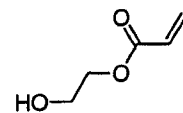
Glycerol dimethacrylate



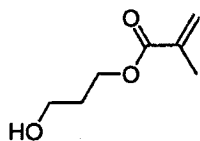
Glycerol diacrylate



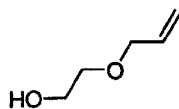
Hydroxy ethylmethacrylate



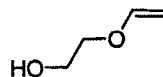
Hydroxy ethylacrylate



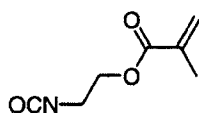
Hydroxy propylmethacrylate



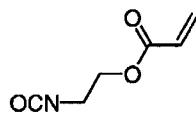
2-Allyloxyethanol



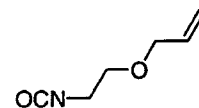
Ethylene glycol vinyl ether



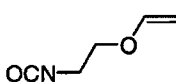
Isocyanato ethyl methacrylate



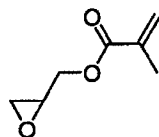
Isocyanato ethyl acrylate



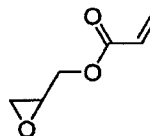
Isocyanato ethyl allyl ether



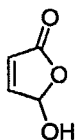
Ethylene glycol vinyl ether



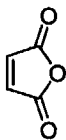
Glycidyl methacrylate



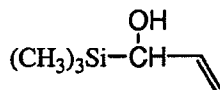
Glycidyl acrylate



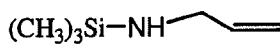
5-hydroxy-2(5H)-furanone



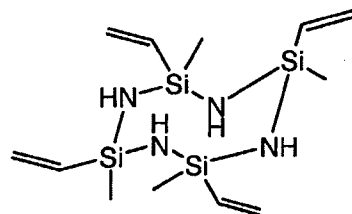
Maleic anhydride



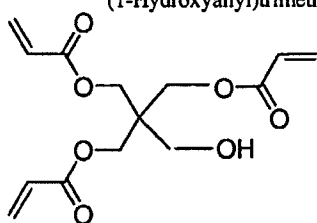
(1-Hydroxyallyl)trimethylsilane



Allylaminotrimethylsilane



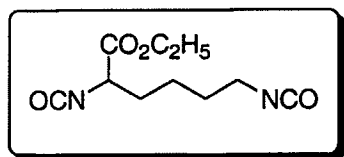
1,3,5,7-Tetravinyl-1,1,3,5,7-tetramethylcyclotetrasilazane



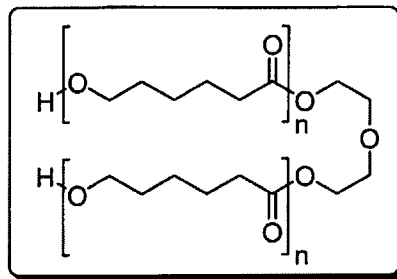
Pentaerythritol triacrylate

Other preferred embodiments are illustrated below. The following discussion is meant by way of example only and should not be considered to limit the invention to the architectures, functional groups or the ways in which the components are bonded and cured which are set out.

- 5 In one preferred embodiment the linker component may be formed from ethyl lysine diisocyanate and poly(caprolactone diol).

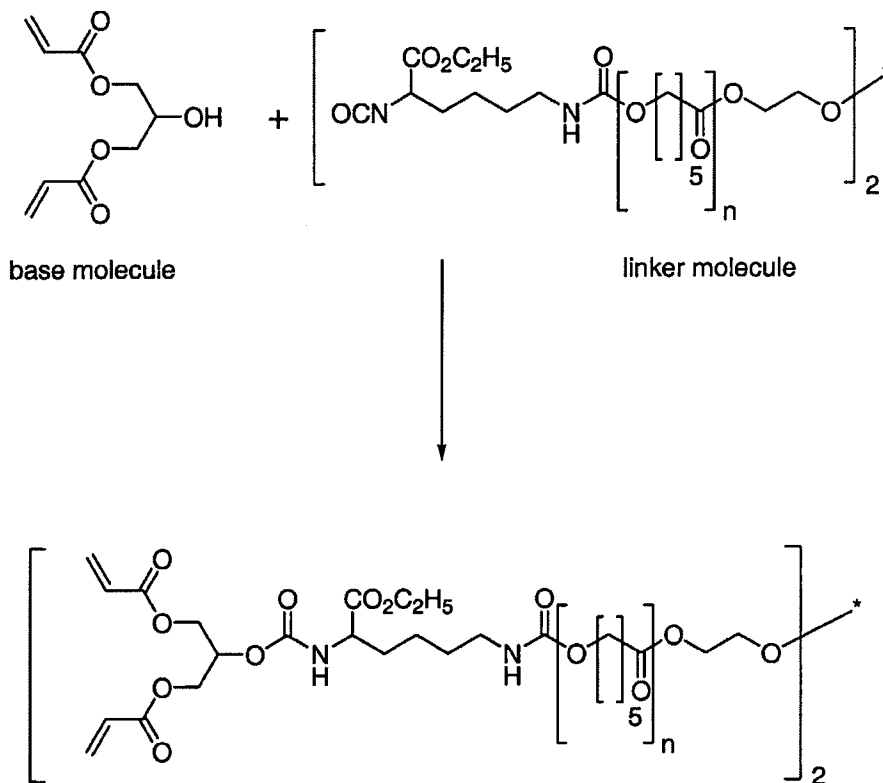


ethyl lysine diisocyanate (ELDI)



poly(caprolactone diol)

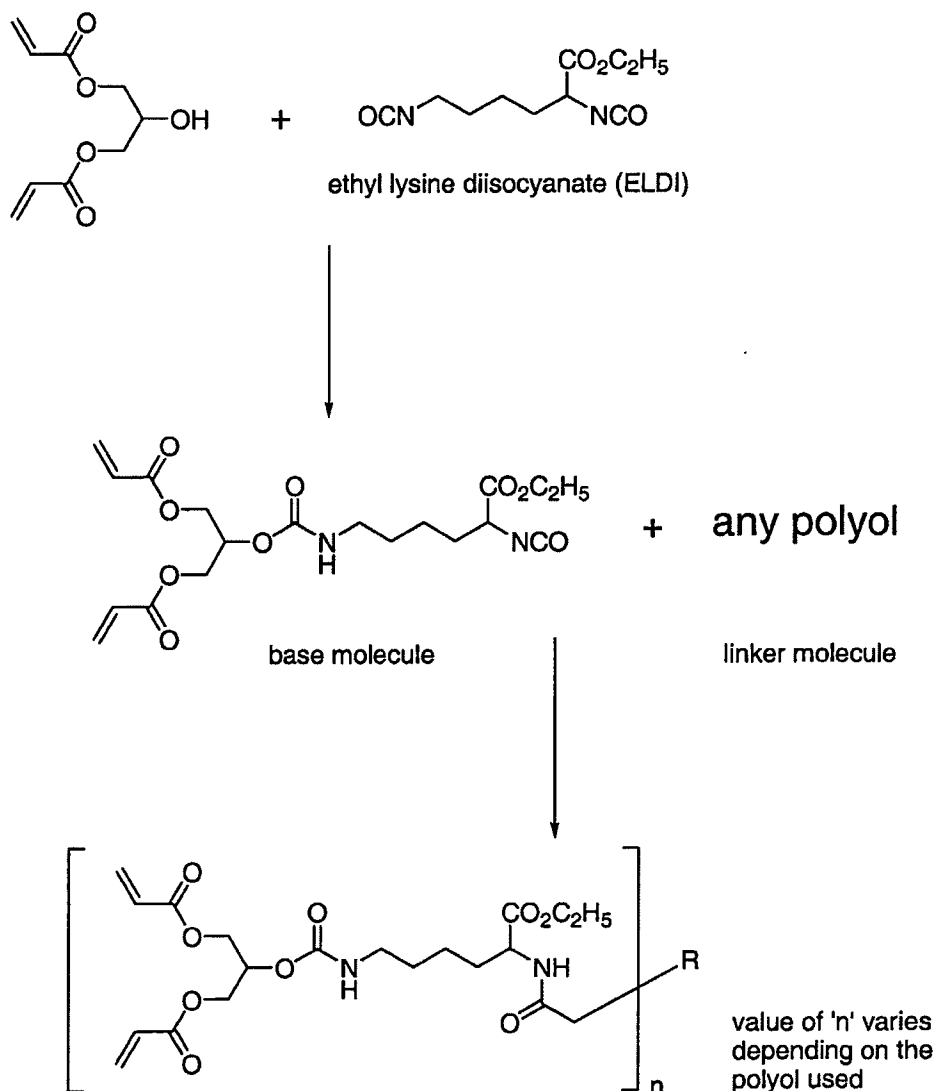
The polymer composition cures in a first stage as set out hereinbelow:



The base molecule in this example is commercially available glycerol dimethyl acrylate which is used to end cap the linker molecule. The linker is easily prepared by mixing appropriate ratios of a polycaprolactone diol of

molecular weight approximately 400, and ethyl lysine diisocyanate, the ratios being selected to determine the properties required for the end product. The hydrophilicity of the end product can be adjusted depending on the desired application by varying the ratio of the acrylate and the polyol. A catalyst in this case is not essential; however, the presence of stannous octoate increases the rate of reaction between the isocyanate and the hydroxyl group. The composition is flowable, injectable through an 18 gauge needle, and can be cured with monochromatic visible light in the presence of suitable radical initiators.

An alternative preferred embodiment is based on a base molecule made from glycerol dimethacrylate and ethyl lysine diisocyanate with the linker being a polyol.

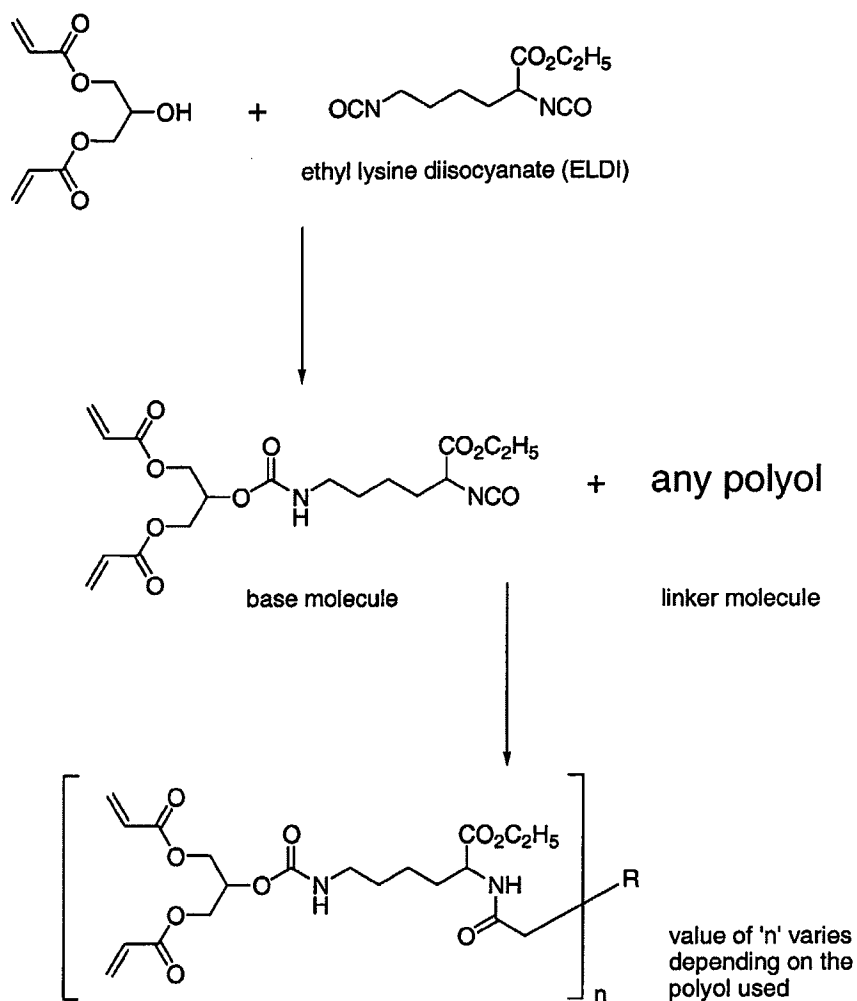


In this case any polyol or an amino functional molecule having an appropriate molecular weight in the range 200 to about 2000 and any architecture, linear or branched, can be used as the linker in the composition.

In the above examples, generally a molar ratio of 4:1 of linker: base molecule is present. Water may be a dispersant in the above preferred embodiments. Preferably up to 40% water by weight may be mixed to achieve a stable emulsion like mixture. In such cases the residual isocyanate groups react with water to produce carbon dioxide and function as a blowing agent to create porosity. Consequent generation of amino groups enhances the hydrophilicity of the surface of the cured polymer. Further increases in hydrophilicity may be achieved by the inclusion of radically polymerizable monomers as additional components in the composition, capable of acting as compatibilizers which alter the hydrophilicity/hydrophobicity ratio thereby adjusting the polymer composition to the environment of use.

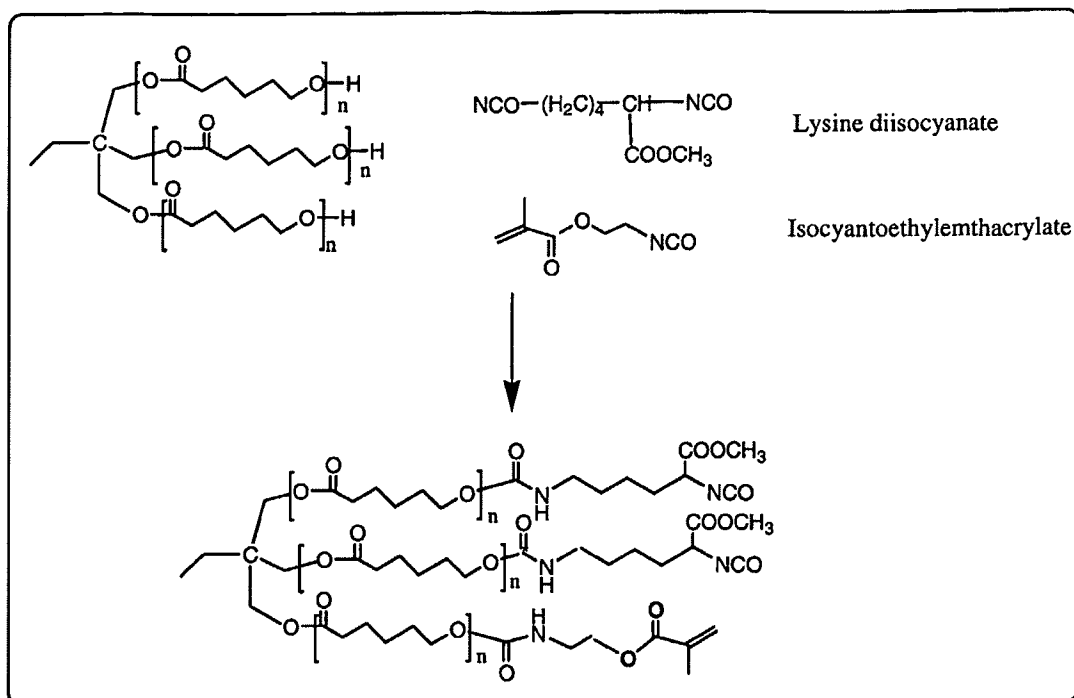
Another alternative preferred embodiment is based on a base molecule made from glycerol dimethacrylate and ethyl lysine diisocyanate with the linker being a polyol.

19



In this case any polyol or an amino functional molecule having an appropriate molecular weight in the range 200 to about 2000 and any
 5 architecture, linear or branched, can be used as the linker in the composition

In another preferred embodiment dual cure polymer compositions may also be prepared using caprolactone triol as linker, and an oligomer of methyl lysine diisocyanate, and isocyanato ethylmethacrylate as base:



The compositions illustrated above can be cured by conventional photolytic or radiation triggered initiators and matching monochromatic light of any radiation frequency. Any initiator compound or compounds which will form initiator radicals so as to effect free radical polymerization are suitable. For the above examples, curing can be achieved with camphorquinone together with a tertiary amine (e.g. dimethyl aminoethyl acrylate), and visible light centred approximately at 450nm. Complete curing can be achieved within very short periods of time, depending on the levels of the initiator. With 0.1% of initiator a 20-60 second burst of blue light at 1000mW/cm² is sufficient for complete cure.

The ratio of base to linker molecule can be varied to alter the properties of the cured material. For example, using a linker that is considered to be soft, such as poly(ε-caprolactone diol) in excess with less cross linking enables the final material to be softer with high elongation. Alternatively increased cross linking allows the preparation of harder material with less elongation.

The following examples are intended to illustrate the scope of the invention and to enable reproduction and comparison. They are not intended to limit the scope of the disclosure in any way.

EXAMPLES

The following examples illustrate the preparation of prepolymer compositions suitable for second and further curing stages.

Examples 1-5 illustrate the preparation of linker molecules for use in the composition of the invention.

Example 1

Materials: Polycaprolactone diol (PCLD 400) with molecular weight Mn 400 (Era Polymers or Aldrich) was pre-dried to eliminate any residual water at 90°C under vacuum (40mm Hg) for at least 2 hours prior to use. Ethyl 2,6-diisocyanato hexanoate (ELDI) (Kyowa Hakko Kogyo Co.Ltd) was used as received from supplier.

Procedure: Pre-dried PCL400 (6.0g, 15 mmol) was weighed in to a round bottom flask equipped with a magnetic stirrer, a nitrogen inlet and a drying tube. ELDI (8.48g, 37.5 mmol) was added to the flask while stirring and heated at 70°C for two hours under nitrogen. The heating bath was removed and the mixture was stirred over night at room temperature. The resultant product was degassed and stored under a nitrogen atmosphere in the freezer. Stannous 2-ethyl hexanoate (0.1% of total mass) may be employed as a catalyst to accelerate the reaction.

Example 2

Materials: Polycaprolactone diol (PCLD1000) with molecular weight Mn 1000 (Era Polymers or Aldrich) was pre-dried to eliminate any residual water at 90°C under vacuum (40mm Hg) for at least 2 hours prior to use. Ethyl 2,6-diisocyanato hexanoate (ELDI) (Kyowa Hakko Kogyo Co.Ltd) was used as received from supplier.

Procedure: PCL1000 (15.00g, 15 mmol) pre-dried as described above was weighed in to a round bottom flask equipped with a magnetic stirrer, a nitrogen inlet and a drying tube. Ethyl 2,6-diisocyanato hexanoate (8.48g, 37.5 mmol) was rapidly added to the diol while with rapid stirring at 40°C and heated at 70°C for two hours. The heating bath was removed and the mixture was stirred over night at room temperature under a nitrogen atmosphere. The mixture thickened on reaction between the diol and the diisocyanate. The mixture was degassed and stored over nitrogen in the freezer. Stannous 2-ethyl hexanoate (0.1% of total mass) may be employed as a catalyst to accelerate the reaction.

Example 3

Materials: Polycaprolactone diol (PCLD2000) with molecular weight Mn 2000 (Era Polymers or Aldrich) was pre-dried to eliminate any residual water at 90°C under vacuum (40mm Hg) for at least 2 hours prior to use. Ethyl 2,6-diisocyanato hexanoate (ELDI) (Kyowa Hakko Kogyo Co.Ltd) was used as received from supplier.

Procedure: PCL2000 (7.50g, 3.75 mmol) pre-dried as described above was weighed in to a round bottom flask equipped with a magnetic stirrer, a nitrogen inlet and a drying tube. Ethyl 2,6-diisocyanato hexanoate (2.11g, 9.38 mmol) was rapidly added to the diol while with rapid stirring at 40°C and heated at 70°C for two hours. The heating bath was removed and the mixture was stirred over night at room temperature under a nitrogen atmosphere. The mixture thickened on reaction between the diol and the diisocyanate. The mixture was degassed and stored over nitrogen in the freezer. At ambient temperature the mixture is a solid, however, melts at 70°C.

Example 4

Materials: Poly(lactic acid) diol (PLA400) with molecular weight Mn 417 was pre-dried to eliminate any residual water at 90°C under vacuum (40mm Hg) for at least 2 hours prior to use. Ethyl 2,6-diisocyanato hexanoate (ELDI) (Kyowa Hakko Kogyo Co.Ltd) was used as received from supplier.

Procedure: PLA400 (3.12g, 7.5 mmol) pre-dried as described above was weighed in to a round bottom flask equipped with a magnetic stirrer, a nitrogen inlet and a drying tube. Ethyl 2,6-diisocyanato hexanoate (4.24g, 18.75 mmol) was rapidly added to the diol while with rapid stirring at 40°C and heated at 70°C for two hours. The heating bath was removed and the mixture was stirred over night at room temperature under a nitrogen atmosphere. The mixture thickened on reaction between the diol and the diisocyanate, was degassed and stored over nitrogen in the freezer. Dibutyl tin octanoate (0.1% of total mass) may be employed as a catalyst to accelerate the reaction.

Example 5

Materials: Polytetramethylene ether glycol (PTMEG1000) with molecular weight Mn 1000 (Aldrich) was pre-dried to eliminate any residual water at 90°C under vacuum (40mm Hg) for at least 2 hours prior to use. 4,4'-

methylenebis(phenyl isocyanate) (MDI) (Aldrich) was used as received from supplier.

Procedure: PTMEG1000 (10.35g, 10.30 mmol) pre-dried as described above was weighed in to a round bottom flask equipped with a magnetic stirrer, a
5 nitrogen inlet and a drying tube. MDI (6.52g, 26 mmol) was rapidly added to the diol while with rapid stirring at 80°C and heated at 80°C for two hours. The heating bath was removed and the mixture was stirred over night at room temperature under a nitrogen atmosphere. The mixture thickened on reaction between the diol and the diisocyanate, was degassed and stored over nitrogen in
10 the freezer. This procedure was adopted from the technical bulletin of "DuPont Terathane", available from the web.

Examples 6-9 illustrate the preparation of prepolymer for use as a base molecule according to the invention.

Example 6

15 Molar ratio; NCO:Acrylate (2:1)

Materials: Polycaprolactone triol (MW 300, Aldrich) was dried under vacuum (0.1 torr) at 80°C over night. Methyl 2,6-diisocyanato hexanoate (MLDI, Kyowa Yuka Co., Ltd, Japan) and Isocyanatoethyl methacrylate and 2,6-di-tert-butyl-4-methylphenol were used as received. All the glassware used was
20 thoroughly cleaned and dried at 105°C overnight in an oven before use.

Procedure: Predried polycaprolactone triol (4.0 g) was weighed in a dry three-neck flask equipped with a magnetic stirrer, nitrogen inlet and drying tube. Methyl 2,6-diisocyanato hexanoate (MLDI) (5.65 g) was then added to the flask followed by isocyanatoethyl methacrylate (IEM) (2.06 g) followed by 2,6-di-tert-butyl-4-methylphenol (0.002g, 0.1 wt% of IEM) under nitrogen. The reaction
25 mixture was stirred and heated in the dark to 70°C for 2h under nitrogen atmosphere. The homogenous polymer mixture was degassed under vacuum (0.1 torr) at 50°C before it was transferred to a vial under nitrogen atmosphere and stored in the refrigerator. The molecular weight and viscosity of the prepolymer were determined by gel permeation chromatography (GPC) and
30 showed number average molecular weight and polydispersity 2116 and 1.21, respectively.

GPC was performed on Water Associates Liquid Chromatograph system (Waters 717) equipped with a differential refractometer and four μ -Styragel

columns (10^5 , 10^4 , 10^3 and 100\AA). The mobile phase was tetrahydrofuran (THF) at a flow rate of 1 mL/min. Prepolymer was dissolved in THF by warming the solution at 50°C for 1h and filtered through 0.45 micron filter before analysis. The system was calibrated with narrow disperse polystyrene standards and molecular weights are reported as polystyrene equivalents. The viscosity was measured using Bohlin Rheometer (CSR10) at 23°C .

Example 7

Molar ratio; NCO: Acrylate (2.5:0.5)

Materials: Polycaprolactone triol (MW 300, Aldrich) was dried under vacuum (0.1 torr) at 80°C over night. Methyl 2,6-diisocyanato hexanoate (MLDI, Kyowa Yuka Co., Ltd, Japan) and isocyanatoethyl methacrylate and 2,6-di-tert-butyl-4-methylphenol were used as received. All the glassware used was thoroughly cleaned and dried at 105°C overnight in an oven before use.

Procedure: Predried polycaprolactone triol (4.0 g) was weighed in a dry three-neck flask equipped with a magnetic stirrer, nitrogen inlet and drying tube. Methyl 2,6-diisocyanato hexanoate (MLDI) (7.07 g) was then added to the flask followed by isocyanatoethyl methacrylate (IEM) (1.03 g) followed by 2,6-di-tert-butyl-4-methylphenol (0.001g, 0.1 wt% of IEM) under nitrogen. The reaction mixture was stirred and heated in the dark to 70°C for 2h under nitrogen atmosphere. The homogenous polymer mixture was degassed under vacuum (0.1 torr) at 50°C before it was transferred to a vial under nitrogen atmosphere and stored in the refrigerator.

Example 8

Preparation of prepolymer with initiator 2-hydroxy-1-[4-(hydroxyethoxy)phenyl]-2-methyl-1-propanone (Irgacure) covalently attached to a base prepolymer.

Molar ratio: NCO:Acrylate: (1.8:1.0)

Materials: Polycaprolactone triol (MW 300, Aldrich) was dried under vacuum (0.1 torr) at 80°C over night. Methyl 2,6-diisocyanato hexanoate (MLDI, Kyowa Yuka Co., Ltd, Japan) and isocyanatoethyl methacrylate and 2,6-di-tert-butyl-4-methylphenol (Aldrich) and Irgacure 2959 (Ciba) were used as received. All the glassware used was thoroughly cleaned and dried at 105°C overnight in an oven before use.

Procedure: Predried polycaprolactone triol (5.0 g) was weighed in a dry three-neck flask equipped with a magnetic stirrer, nitrogen inlet and drying tube. Methyl 2,6-diisocyanato hexanoate (MLDI) (6.36 g) was then added to the flask followed by isocyanatoethyl methacrylate (IEM) (2.56 g) followed by 2,6-di-tert-butyl-4-methylphenol (0.002g, 0.1 wt% of IEM) and Irgacure2959 (0.786g) under nitrogen. The reaction mixture was stirred and heated in the dark to 70°C for 2-2.5h under nitrogen atmosphere. The homogenous polymer mixture was degassed under vacuum (0.1 torr) at 50°C before it was transferred to a vial under nitrogen atmosphere and stored in the refrigerator.

10 Example 9

Preparation of a Glycerol based prepolymer NCO: Acrylate (1:3)

Materials: Glycerol (Aldrich) was dried under vacuum (0.1 torr) at 80°C over night. Methyl 2,6-diisocyanato hexanoate (MLDI, Kyowa Yuka Co., Ltd, Japan) and Isocyanatoethyl methacrylate and 2,6-di-tert-butyl-4-methylphenol were used as received. All the glassware used was thoroughly cleaned and dried at 105°C overnight in an oven before use.

Procedure: Glycerol (2.50 g) was weighed in a dry three-neck flask equipped with a magnetic stirrer, nitrogen inlet and drying tube. Isocyanatoethyl methacrylate (IEM) (12.6 g) was then added to the flask followed by 2,6-di-tert-butyl-4-methylphenol (0.0016g, 0.1 wt% of IEM) under nitrogen. The reaction mixture was stirred and heated in dark to 70°C for 22h under nitrogen atmosphere. The homogenous mixture was degassed under vacuum (0.1 torr) at 70°C before it was transferred to a vial under nitrogen atmosphere and stored in the refrigerator. The number average molecular weight and polydispersity of the prepolymer were 607 and 1.02, respectively based on GPC analysis.

Examples 10 to 17, except example 15, illustrate curing of prepolymer using either visible or UV light. Following is a brief description of the procedure used.

For visible light curing the light source used was supplied by 3M industries. The model used was ESPE Elipar Freelight 2, which emits in the range of 430-480nm and at an intensity of 1000 mW/cm². The experiments conducted in Teflon wells, curing from one side.

Procedure: The degassed pre-polymer was weighed in to a glass sample vial. Camphorquinone (0.1%) and N,N dimethyl amino ethyl methacrylate

(DMAEMA) (0.2%) (or any other sensitizer) was added, mixed with the prepolymer. Water (with or without gelatin beads) is added (if desired) up to 40% of the mass, mixed to a creamy consistency and the mixture taken up in a syringe. The composition with camphorquinone needs to be kept in a dark environment as
5 natural light triggers the curing process. The polymer mixture is poured in to a cylindrical Teflon cavity and irradiated using the blue lamp for 1-2 minutes, during which time the curing occurs. The solid plug can be removed immediately from the mould as 1-2 minutes curing turns it in to a hard solid.

Example 10

10 Degassed prepolymer (1.00 g) of Polycaprolactone triol with MLDI prepared in Example 1 was weighed in to a cavity (20x20x10 mm) made in a Teflon block. Degassed and dried polycaprolactone triol of molecular weight (MW 300, 0.1819 g), camphorquinone (0.1 wt% 0.0018g) and N,N-dimethyl amino ethyl methacrylate (DMAEMA) (0.003g) was mixed together and added to the
15 prepolymer. The mixture was stirred for several minutes in dark and then gelatin beads (10-50 % by weight) was added and stirred further for 5 min. The mixture which was a viscous and injectable liquid, was taken into a 2.5 ml syringe and dispensed 0.29 g into each cylindrical cavity (6 mm D x 12 mm L) in a multi-cavity Teflon mould, sealed and cured for 10 sec with blue light (430-460 nm) to give
20 porous cylindrical polymer test specimens.

The following examples illustrate UV (photo)curing.

Example 11

Tagged UV initiator (Irgacure 2959)

25 Degassed prepolymer (0.35 g) of PCLT and IEM and MLDI prepared in Example 8 was weighed in a polypropylene mould. N,N-dimethyl amino ethyl methacrylate (DMAEMA) (0.015 g) and gelatine beads (0.1g) was added to the prepolymer. The mixture was mixed well for few minutes and then cured with UV light (365 nm, 1 MW/cm²@8cm, Spectraline) to give a porous polymer test specimens.

30 Example 12

Glycerol dimethacrylate (91 mg, 0.40 mmolL) was mixed with the composition made in example 5(100 mg, 0.10 mmolL) and 1mg of stannous-2-ethylhexanoate in a glass vial. The mixture was allowed to stand under nitrogen for 4 hours in the dark and CIBA UV initiator Darocur (5 mg) added. The mixture

was placed under a UV lamp (365 nm, 1 MW/cm²@8cm, Spectraline) for 1 hour at which time the mixture hardened to a solid, with good adhesion to glass. A control experiment without a radical initiator remained non-cross linked.

Example 13

5 Glycerol dimethacrylate (45.60 mg, 0.20 mmol) was mixed with the composition of example 1 (170 mg, 0.20 mmol) and 0.5mg of CIBA UV initiator Darocur in a glass vial. The mixture was allowed to stand under nitrogen for 4 hours in the dark and exposed to UV light (365 nm, 1 MW/cm²@8cm, Spectraline) for 1 hour at which time the mixture cured to a hard solid, with good
10 adhesion to glass. A control experiment without a radical initiator remained non-cross linked.

The following example illustrates the use of water to create a porous material.

Example 14

15 Glycerol dimethacrylate (45.60 mg, 0.20 mmol) was mixed with a prepolymer according to example 1 (170 mg, 0.20 mmol) and 0.5mg of CIBA UV initiator Darocur in a glass vial. The mixture was allowed to stand under nitrogen for 4 hours in the dark and 100mg of water added. The mixture was mixed to a creamy composition and exposed to UV light (365 nm, 1 MW/cm²@8cm, Spectraline) for 1 hour at which time the mixture cured to a hard porous solid. A
20 control experiment without a radical initiator remained non-cross linked.

The following example illustrates the use of a redox initiator in the second curing step.

Example 15

25 Degassed prepolymer (0.02 g) prepared in Example 9 was weighed in to a cavity (20x20x10 mm) made in a Teflon block. A solution of Ammonium per sulfate (0.04g/ml) (20 micro litre) and Sodium metabisulfite (0.04g/ml) (20 micro litre) and N,N,N,N-tetramethylethylenediamine (sensitizer) 0.1 wt% were added to the prepolymer. The mixture was stirred for several minutes and dispensed 0.29 g
30 into each cylindrical cavity (6 mm D x 12 mm L) in a cavity Teflon mould covered with glass plate and cured at 37°C to give porous cylindrical polymer test specimens.

Example 16

Degassed prepolymer (1.00 g) of Polycaprolactone triol with MLDI prepared in Example 6 was weighed in to a cavity (20x20x10 mm) made in a Teflon block. Degassed and dried polycaprolactone triol of molecular weight (MW 5 300, 0.1819 g), initiator camphorquinone (0.1 wt% 0.0018g) and N,N-dimethyl amino ethyl methacrylate (DMAEMA) (0.003g) was mixed together and added to the prepolymer. The mixture was stirred for several minutes in the dark and then gelatine beads (10-50 % by weight) was added and stirred further for 5 min. The mixture which was a viscous and injectable liquid, was taken into a 2.5 ml syringe 10 and dispensed 0.29 g into each cylindrical cavity (6 mm D x 12 mm L) in a multi-cavity Teflon mould, sealed and cured for 10 sec with blue light (430-460 nm) to give porous cylindrical polymer test specimens.

The prepolymer prepared above was also cured with UV light (365 nm) using UV initiator Irgacure 2959 to give a porous polymer test specimens.

15 Example 17

ELDI (40.14g) was added to four arm poly(D/L-lactic acid) (39.47g, Mn 889) in a round bottom flask. Stannous 2-ethyl hexanoate (40mg, 0.05%) was added and the mixture allowed to stir for 12 hours.

Part of the above reaction mixture (24.96g) was placed in a round bottom 20 flask, pentaerythritol triacrylate (16.6g) and MEHQ (32.5mg) added. The mixture (prepolymer) was allowed to stir for 12 hours before use.

No filler: The pre-polymer (6.040g) was weighed in to small beaker and mixed to a creamy consistency before CQ (20mg) and DMAEMA (40mg) was added. The mixtures were degassed for a brief period under moderate vacuum 25 (70mmHg) to remove any air bubbles and samples placed in 6mm x 12mm well plates made of Teflon. Two glass plates were placed on either side of the Teflon block, clamped and the polymer samples in the wells irradiated with blue light for 2-3 minutes. Upon completion of irradiation the samples were left in the mould for 24 hours. Removed the samples from the mould and further cured using blue 30 light to ensure complete curing. The mechanical properties of the resultant structures are described in Table 3.

TCP-20: The pre-polymer (5.478g) and β -tricalcium phosphate (TCP) (1.096g) was weighed in to small beaker and mixed to a creamy consistency

before CQ (19.7mg) and DMAEMA (40mg) was added. The mixtures were degassed for a brief period under moderate vacuum (70mmHg) to remove any air bubbles and samples placed in 6mm x 12mm well plates made of Teflon. Two glass plates were placed on either side of the Teflon block, clamped and the
5 polymer samples in the wells irradiated with blue light for 2-3 minutes. Upon completion of irradiation the samples were left in the mould for 24 hours. Removed the samples from the mould and further cured using blue light to ensure complete curing. The sample specimens were conditioned at ambient temperature for 10 days before testing for mechanical properties, the details of
10 which are provided in Table 3.

HA-20: The pre-polymer (5.731g) and hydroxy apatite (1.146g) was weighed in to small beaker and mixed to a creamy consistency before CQ (20mg) and DMAEMA (40mg) was added. The mixtures were degassed for a brief period under moderate vacuum (70mmHg) to remove any air bubbles and samples
15 placed in 6mm x 12mm well plates made of Teflon. Two glass plates were placed on either side of the Teflon block, clamped and the polymer samples in the wells irradiated with blue light for 2-3 minutes. Upon completion of irradiation the samples were left in the mould for 24 hours. Removed the samples from the mould and further cured using blue light to ensure complete curing. The
20 mechanical properties of the resultant structure are described in Table 3.

Example 18

ELDI (20.85g) was added to four arm poly(D/L-lactic acid) (20.85g, Mn 889) in a round bottom flask. The mixture allowed to stir for 12 hours. Part of the above reaction mixture (10.76g) was placed in a round bottom flask,
25 Glycerol diacrylate (5.47g) and BHT (38mg) added. The mixture was allowed to stir for 12 hours before use.

No filler: The pre-polymer (4.40g) was weighed in to small beaker and mixed to a creamy consistency before CQ (4.4mg) and DMAEMA (20mg) was added. The mixtures were degassed for a brief period under moderate vacuum
30 (70mmHg) to remove any air bubbles and samples placed in 6mm x 12mm well plates made of Teflon. Two glass plates were placed on either side of the Teflon block, clamped and the polymer samples in the wells irradiated with blue light for 2-3 minutes. Upon completion of irradiation the samples were left in the mould for 24 hours. Removed the samples from the mould and further cured using blue

light to ensure complete curing. The mechanical properties of the resultant sample are described in Table 3.

HA-10: The pre-polymer (4.40g) and hydroxy apatite (0.44g) was weighed in to small beaker and mixed to a creamy consistency before CQ (5mg) and DMAEMA (10mg) was added. The mixtures were degassed for a brief period under moderate vacuum (70mmHg) to remove any air bubbles and samples placed in 6mm x 12mm well plates made of Teflon. Two glass plates were placed on either side of the Teflon block, clamped and the polymer samples in the wells irradiated with blue light for 2-3 minutes. Upon completion of irradiation the samples were left in the mould for 24 hours. Removed the samples from the mould and further cured using blue light to ensure complete curing. The mechanical properties of the resultant samples are described in Table 3.

Example 19

ELDI (4.52g) was added to a solution of four arm -poly(D/L-lactic acid-co-glycolic acid) (20g, Mn 4000) in dichloromethane (20mL). The mixture allowed to stir for 12 hours.

Glycerol diacrylate (4.56g) in dichloromethane (5mL) and BHT (35mg) added. The mixture was allowed to stir for 12 hours and the solvent removed under high vacuum before use.

No filler: The pre-polymer (3.34g) was weighed in to small beaker and mixed to a creamy consistency before CQ (10mg) and DMAEMA (100mg) was added. The mixtures were degassed for a brief period under moderate vacuum (70mmHg) to remove any air bubbles and samples placed in 6mm x 12mm well plates made of Teflon. Two glass plates were placed on either side of the Teflon block, clamped and the polymer samples in the wells irradiated with blue light for 2-3 minutes. Upon completion of irradiation the samples were left in the mould for 24 hours. Removed the samples from the mould and further cured using blue light to ensure complete curing. The mechanical properties of the resultant structures are described in Table 3.

Example 20

ELDI (4.70g) was added to a solution of four arm -poly(D/L-lactic acid-co-glycolic acid, 3:1) (6.18g, Mn 1228) in dichloromethane (2mL). The mixture allowed to stir for 12 hours.

Pentaerythritol triacrylate (5.98g) and BHT (6mg) added. The mixture was allowed to stir for 12 hours and the solvent reduced under high vacuum before use.

TCP-20: The pre-polymer (5.079g) and β -tricalcium phosphate (TCP) (1.015g) was weighed in to small beaker and mixed to a creamy consistency before CQ (~15mg) and DMAEMA (~30mg) was added. The mixtures were degassed for a brief period under moderate vacuum (70mmHg) to remove any air bubbles and samples placed in 6mm x 12mm well plates made of Teflon. Two glass plates were placed on either side of the Teflon block, clamped and the polymer samples in the wells irradiated with blue light for 2-3 minutes. Upon completion of irradiation the samples were left in the mould for 24 hours. Removed the samples from the mould and further cured using blue light to ensure complete curing. The sample specimens were conditioned at ambient temperature for 7 days before testing for mechanical properties. The mechanical properties are described in Table 3.

Example 21

ELDI (4.74g) was added to a solution of four arm -poly(D/L-lactic acid-co-glycolic acid) (6.576g, Mn 1228) in dichloromethane (2mL). The mixture allowed to stir for 12 hours.

Pentaerythritol triacrylate (6.20g) and BHT (6mg) added. The mixture was allowed to stir for 12 hours and the solvent reduced under high vacuum before use.

TCP-20: The pre-polymer (5.043g) and β -tricalcium phosphate (TCP) (1.008g) was weighed in to small beaker and mixed to a creamy consistency before CQ (~15mg) and DMAEMA (~30mg) was added. The mixtures were degassed for a brief period under moderate vacuum (70mmHg) to remove any air bubbles and samples placed in 6mm x 12mm well plates made of Teflon. Two glass plates were placed on either side of the Teflon block, clamped and the polymer samples in the wells irradiated with blue light for 2-3 minutes. Upon completion of irradiation the samples were left in the mould for 24 hours. Removed the samples from the mould and further cured using blue light to ensure complete curing. The sample specimens were conditioned at ambient temperature for 7 days before testing for mechanical properties. The mechanical properties are described in Table 3.

Table 3: Mechanical properties of polymer compositions prepared according to examples 17-21.

Sample code	Ultimate Compression Strength (MPa)	Compression Strength @ Yield Point (MPa)	Compression Modulus (GPa)	Formulation/comments
Example 17 No Filler	60±10	32±1	0.9±0.2	PLA (900), PETA/No filler
Example 17 TCP-20	65±10	55±10	2.1±0.5	PLA (900), PETA, TCP (20%)
Example 17 HA-20	61 ±9	32±5	1.1±0.1	PLA (900), PETA, HA (20%)
Example 18 No Filler	38±10	33±7	1.1±0.3	PLA (900), GDA, no filler
Example 18 HA-10	44±8	43±6	1.3±0.3	PLA(900), GDA, HA (10%)
Example 19 No Filler	-	59±3	26±2 (MPa)	P[LA-GA, 3:1](4000), GDA
Example 20 TCP-20	83±10	32±8	1.8±0.3	P(LA:GA, 3:1] (1228); PETA, TCP(20%)
Example 21 TCP-20	76.0±0.2	32±6	1.6±0.1	P(LA:GA, 1:1] (1228); PETA, TCP (20%)

5 Example 22

Preparation of polymer: ELDI (12.43g) was added to linear poly(glycolic acid) (10.3g, Mn 411) in a round bottom flask. Stannous 2-ethyl hexanoate (22mg, 0.1%) was added and the mixture allowed to stir for 24 hours.

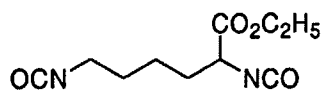
- Part of the above reaction mixture (10.37g) was placed in a beaker,
- 10 Glycerol diacrylate (1.37g), camphoquinone (23.5mg, 0.2%), DMAEMA (47mg, 0.4%) an gelatine beads dispersed in water (2.4g, 20%) added. and BHT (38mg) added. The compositin was mixed thoroughly before cured. Samples were placed in 6mm (D) x 12mm (H) well plates made of Teflon, glass plates placed on either side, clamped and the samples irradiated with blue light for 2-3 minutes.
- 15 Upon completion of irradiation the samples were left in the mould for 24 hours.

The samples were removed from the mould and further cured using blue light to ensure complete curing.

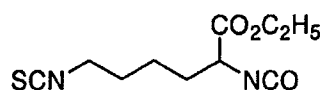
Example 23

The use of sulphur analogs of methyl or ethyl lysine diisocyanate in place of or in addition to any other isocyanates described herein is expected to result in polymer compositions in accordance with the examples described above.

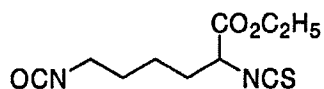
5 Possible analogs are:



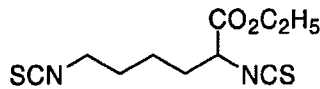
ethyl lysine diisocyanate (ELDI)



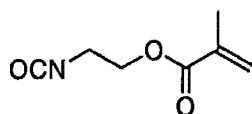
ethyl 2-thio-6-diisocyanato hexanoate



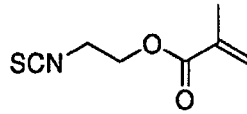
ethyl 6-thio-2-diisocyanato hexanoate



ethyl lysine dithio diisocyanate



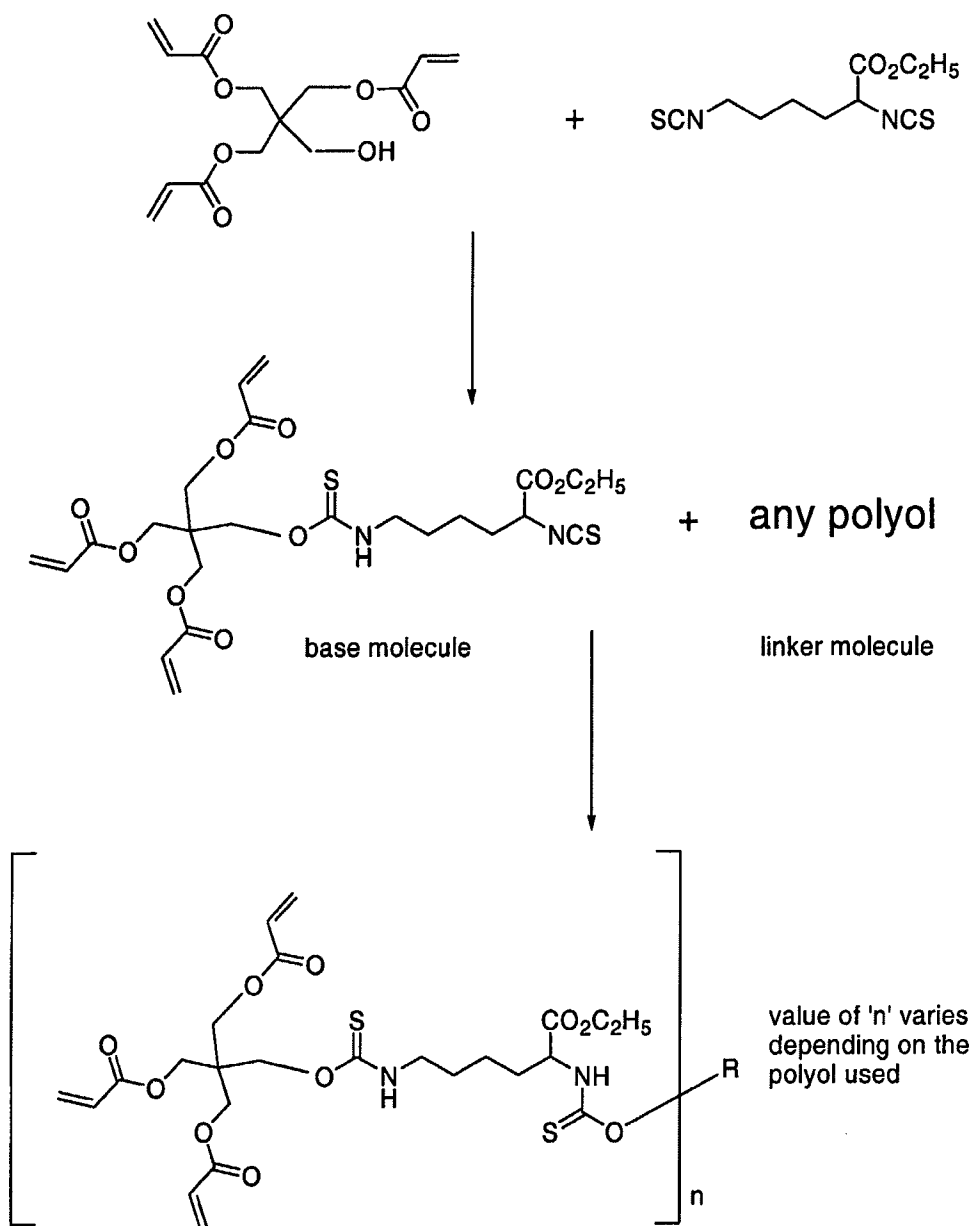
isocyanato ethyl methacrylate (IEM)



thioisocyanato ethyl methacrylate (IEM)

For example, pentaerythritol triacrylate when added to ethyl lysine dithio diisocyanate, is expected to result in the thio variant of the base molecule below. Upon addition of this base molecule to any linker polyol, the link is expected to result in thiourethane functionality.

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The compositions of the invention in their most preferred embodiment use free radical polymerisation of ethylenically unsaturated end groups attached to the prepolymers to form a cross linked network. Porus and non-porus polymers have been formed following the application of electromagnetic radiation in both visible (blue light ~ 440nm) and UV regions, for short periods. The non-cross linked prepolymers prepared are low molecular weight liquids, injectable through an 18 gauge needle and may be cross linked on demand under wet conditions. The prepolymers designed may be biodegradable.

It will be appreciated that the invention is not limited to the examples which are merely illustrative. Other polymer compositions not specifically exemplified

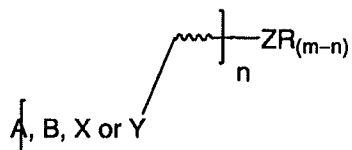
but falling within the spirit of the invention are considered within the scope of the invention.

CLAIMS:

1. A polymer composition for use at least in surface coating applications and in fabrication of rigid foams with load bearing capacity comprising a base molecule, a linker molecule and at least one initiator compound, said base
5 molecule having at least two differing functionalities, and said linker molecule having a functionality reactive with at least one of said functionalities of said base molecule, the first of said at least two functionalities of said base molecule enabling a first curing stage of said polymer composition by reaction with the functionality of said linker molecule, and the second and any further functionality
10 of said base molecule enabling second and optionally further curing stages of said polymer composition, said first, second and any further curing stages being capable of activation simultaneously or independently of each other as required.
2. A polymer composition as claimed in Claim 1 which is biodegradable.
3. A polymer composition as claimed in Claim 1 which is biostable.
- 15 4. A polymer composition as claimed in any one of Claims 1-3 wherein the base molecule and the linker molecule are, independently of each other, a single organic molecule.
5. A polymer composition as claimed in any one of Claims 1-3 wherein the base molecule and the linker molecule are, independently of each other, an
20 oligomer formed from two or more substrate monomers.
6. A polymer composition as claimed in Claim 5 wherein the base molecule and linker molecule independently of each other, have a molecular weight of less than 2000, preferably less than 1000 and more preferably less than 500.
7. A polymer composition as claimed in any one of Claims 1-6 wherein the
25 base molecule is of the formula 1:

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(a)



$n = 1 \text{ to } m$
 $m = \text{valency of } Z$

I

wherein:

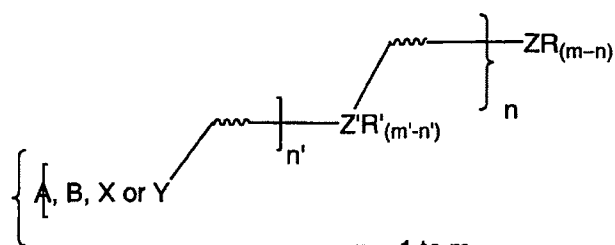
- 5 \sim is one or more repeating units, N & M, are integers, A and B are unsaturated moieties, Z is H, C, O, N, Si or S and the remaining variables are defined as follows:

Atom (Z)	Valancy (m)	n (arms)	R	ZR(m-n)
C	4	1	H, CH ₃ , C ₂ H ₅ or OR	CH ₃ , C(CH ₃) ₃ , C(C ₂ H ₅) ₃ , C(OR) ₃
		2	H, CH ₃ , C ₂ H ₅ or OR	CH ₂ , C(CH ₃) ₂ , C(C ₂ H ₅) ₂ , C(OR) ₂
		3	H, CH ₃ , C ₂ H ₅ or OR	CH, C(CH ₃), C(C ₂ H ₅), C(OR)
		4	none	C (4 arms)
O	2	1	H, CH ₃ , C ₂ H ₅ or OR	OH, O(CH ₃), O(C ₂ H ₅), R cannot be OR
		2	none	O
Si	4	1	H, CH ₃ , C ₂ H ₅ or OR	SiH ₃ , Si(CH ₃) ₃ , Si(C ₂ H ₅) ₃ , Si(OR) ₃
		2	H, CH ₃ , C ₂ H ₅ or OR	SiH ₂ , Si(CH ₃) ₂ , Si(C ₂ H ₅) ₂ , Si(OR) ₂
		3	H, CH ₃ , C ₂ H ₅ or OR	SiH, Si(CH ₃), Si(C ₂ H ₅), Si(OR)
		4	none	Si (4 arms)
H	1	1	none	H (R=0)

8. A polymer composition as claimed in any one of claims 1-6 wherein the base molecule is of the formula II:

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(b)



II
 $n = 1$ to m
 m = valancy of Z
 $n' = 1$ to $(m'-1)$
 m' = valancy of Z'

wherein:

5 \sim is one or more repeating units, N , M , N^1 and M^1 are integers, A and B are unsaturated moieties, Z is H , C , O , N , Si , S and Z^1 is H , C , O , N , Si , S and the remaining variables are defined as follows:

Atom (Z)	Valancy (m)	n (arms)	R	ZR(m-n)
C	4	1	H, CH ₃ , C ₂ H ₅ or OR	CH ₃ , C(CH ₃) ₃ , C(C ₂ H ₅) ₃ , C(OR) ₃
		2	H, CH ₃ , C ₂ H ₅ or OR	CH ₂ , C(CH ₃) ₂ , C(C ₂ H ₅) ₂ , C(OR) ₂
		3	H, CH ₃ , C ₂ H ₅ or OR	CH, C(CH ₃), C(C ₂ H ₅), C(OR)
		4	none	C (4 arms)
O	2	1	H, CH ₃ , C ₂ H ₅ or OR	OH, O(CH ₃), O(C ₂ H ₅), R cannot be OR
		2	none	O
Si	4	1	H, CH ₃ , C ₂ H ₅ or OR	SiH ₃ , Si(CH ₃) ₃ , Si(C ₂ H ₅) ₃ , Si(OR) ₃
		2	H, CH ₃ , C ₂ H ₅ or OR	SiH ₂ , Si(CH ₃) ₂ , Si(C ₂ H ₅) ₂ , Si(OR) ₂
		3	H, CH ₃ , C ₂ H ₅ or OR	SiH, Si(CH ₃), Si(C ₂ H ₅), Si(OR)
		4	none	Si (4 arms)
H	1	1	none	H (R=0)

Atom (Z')	Valancy (m')	n' (arms)	R'	Z'R'(m'-n')
C	4	1	H, CH ₃ , C ₂ H ₅ or OR	CH ₂ , C(CH ₃) ₂ , C(C ₂ H ₅) ₂ , C(OR) ₂
		2	H, CH ₃ , C ₂ H ₅ or OR	CH, C(CH ₃), C(C ₂ H ₅), C(OR)
		3	none	C (4 arms)
O	2	1	none	O
Si	4	1	H, CH ₃ , C ₂ H ₅ or OR	SiH ₂ , Si(CH ₃) ₂ , Si(C ₂ H ₅) ₂ , Si(OR) ₂
		2	H, CH ₃ , C ₂ H ₅ or OR	SiH, Si(CH ₃), Si(C ₂ H ₅), Si(OR)
		3	none	Si (4 arms)
H	1	0	cannot have a group or atom	

9. A polymer composition as claimed in any one of Claims 1-8 additionally comprising one or more of the group consisting of radical inhibitor, sensitizer, promoter, dispersant, porogen, catalyst, pigment and surfactant.

10. A prepolymer composition for use at least in surface coating and adhesive applications and in the fabrication of rigid foams with load bearing capacity, comprising the reaction product of a base molecule, a linker molecule and at least one initiator compound, said base molecule having at least two differing functionalities, and said linker molecule having a functionality reactive with at least one of said functionalities of said base molecule, said reaction product being the result of a first curing stage wherein the first of said at least two functionalities of said base molecule reacts with the functionality of said linker molecule to form said prepolymer composition.

11. A cured polymeric end product for use at least in surface coating and adhesive applications and in the fabrication of rigid foams with load bearing capacity, comprising the reaction product of a base molecule, a linker molecule and at least one initiator compound, said base molecule having at least two differing functionalities, and said linker molecule having a functionality reactive with at least one of said functionalities of said base molecule, said end product being the result of a first curing stage wherein the first of said at least two functionalities of said base molecule reacts with the functionality of said linker molecule and a second curing stage and optionally further curing stages wherein

said initiator compound is activated to affect free radical polymerisation of at least said second functionality of said base molecule.

12. Use of a polymer composition according to any one of Claims 1-9, a prepolymer composition according to Claim 10 or cured end product according to
5 Claim 11 in the preparation of a surface coating.

13. Use of a polymer composition according to any of Claims 1-9, a prepolymer composition according to Claim 10 or cured end product according to Claim 11 in the preparation of a rigid foam with load bearing capacity.

14. Use of a polymer composition according to any one of Claims 1-9, a
10 prepolymer composition according to Claim 10 or cured end product according to Claim 11 in the preparation of an adhesive composition.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU2005/000306

A. CLASSIFICATION OF SUBJECT MATTER												
Int. Cl. ⁷ : C08G 18/10; C09D 167/07, 175/16; C09J 167/07, 175/16												
According to International Patent Classification (IPC) or to both national classification and IPC												
B. FIELDS SEARCHED												
Minimum documentation searched (classification system followed by classification symbols)												
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched												
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) WPAT and JAPIO:- IPC as above and keywords:- dual cure, isocyanate, polyol, polycaprolactone, acrylate, polyurethane, prepolymer and like terms												
C. DOCUMENTS CONSIDERED TO BE RELEVANT												
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.										
X	WO 2002/010292A (PPG INDUSTRIES OHIO, INC.) 7 February 2002 Page 77	1-12										
X	US 4451523A (Nativi et al.) 29 May 1984 Example 1	1-12										
X	US 4424252 A (Nativi) 3 January 1984 Example 1; columns 8-9	1-12										
X	US 4035274 A (McGinniss) 12 July 1977 Example 2	1-12										
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C <input checked="" type="checkbox"/> See patent family annex												
<p>* Special categories of cited documents:</p> <table border="0"> <tr> <td>"A" document defining the general state of the art which is not considered to be of particular relevance</td> <td>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</td> </tr> <tr> <td>"E" earlier application or patent but published on or after the international filing date</td> <td>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</td> </tr> <tr> <td>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</td> <td>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</td> </tr> <tr> <td>"O" document referring to an oral disclosure, use, exhibition or other means</td> <td>"&" document member of the same patent family</td> </tr> <tr> <td>"P" document published prior to the international filing date but later than the priority date claimed</td> <td></td> </tr> </table>			"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention	"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone	"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art	"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family	"P" document published prior to the international filing date but later than the priority date claimed	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention											
"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone											
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art											
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family											
"P" document published prior to the international filing date but later than the priority date claimed												
Date of the actual completion of the international search 1 April 2005		Date of mailing of the international search report 07 APR 2005										
Name and mailing address of the ISA/AU AUSTRALIAN PATENT OFFICE PO BOX 200, WODEN ACT 2606, AUSTRALIA E-mail address: pct@ipaustalia.gov.au Facsimile No. (02) 6285 3929		Authorized officer N.L. KING Telephone No : (02) 6283 2150										

INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU2005/000306

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2003/0153673 A1 (Schwalm et al.) 14 August 2003 Examples 1 and 2	1-12
X	DEMARTEAU, W. et al. Dual UV/Thermally Curable Formulations In Radcure '84 Conference Proceedings, pages 1-1 to 1-10 Entire Document	1-12
X	ICHINOSE, E. et al. Study of Hybrid Curing Behaviour of Radiation Curing Systems In Radtech '91 Conference Proceedings, pages 738 to 752 Entire Document	1-12
X	US 6124370 A (Walton et al.) 26 September 2000 Formulations 6-7	1-11, 13
X	US 5276068 A (Waknine) 4 January 1994 Column 9 lines 37-65; Examples 1-23	1-11, 14

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/AU2005/000306

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report				Patent Family Member			
WO	0210292	AU	63930/00	BR	0017309	CA	2417257
		EP	1305374	MX	PA03000930		
US	4451523	AU	19822/83	CA	1236240	CA	1238443
		EP	0112005	EP	0231548	JP	1163274
		JP	59135263	US	4415604		
US	4424252	AU	19821/83	CA	1238134	EP	0110540
		HK	55588	JP	59102965		
US	4035274	CA	1073142	US	4035273	US	4070258
		US	4130522	US	4166017		
US	2003153673	AU	81894/01	DE	10031258	EP	1294788
		WO	0200754				
US	5276068	JP	62226907	US	5444104	DE	3632868
		CH	674308	DE	3610808		
Due to data integration issues this family listing may not include 10 digit Australian applications filed since May 2001.							
END OF ANNEX							